Table 1. Level 1 Component and Composition Changes for Immediate Release Oral Solid Dosage Forms

Excipient Percent Excipient (w/w) Out of To	
	Dosage Form Weight
Filler	+/- 5%
Disintegrant	
starch	+/- 3%
other	+/- 1%
Binder	+/- 0.5%
Lubricant	
calcium or magnesium stearate	+/- 0.25%
other	+/- 1%
Glidant	
talc	+/- 1%
other	+/- 0.1%
Film coat	+/- 1%

Table 2. Level 2 Component and Composition Changes for Immediate Release Oral Solid Dosage Forms

TORIE			
Excipient	Percent Excipient (w/w) Out of Total Target		
	Dosage Form Weight		
Filler	+/- 10%		
Disintegrant			
starch	+/- 6%		
other	+/- 2%		
Binder	+/- 1%		
Lubricant			
calcium or magnesium stearate	+/- 0.5%		
other	+/- 2%		
Glidant			
talc	+/- 2%		
other	+/- 0.2%		
Film coat	+/- 2%		

Figure 1

Table 3. Level 3 Component and Composition Changes for Immediate Release Oral Solid Dosage Forms

Excipient	Percent Excipient (w/w) Out of Total Target	
	Dosage Form Weight	
Filler	Greater than +/- 10%	
Disintegrant		
starch	Greater than +/- 6%	
other	Greater than +/- 2%	
Binder	Greater than +/- 1%	
Lubricant		
calcium or magnesium stearate	Greater than +/- 0.5%	
other	Greater than +/- 2%	
Glidant		
talc	Greater than +/- 2%	
other	Greater than +/- 0.2%	
Film coat	Greater than +/- 2%	

Table 4. Level 1 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (nonrelease controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Target
	Dosage Form Weight
Filler	+/- 5%
Disintegrant	
starch	+/- 3%
other	+/- 1%
Binder	+/- 0.5%
Lubricant	
calcium or magnesium stearate	+/- 0.25%
other	+/- 1%
Glidant	
talc	+/- 1%
other	+/- 0.1%
Film coat	+/- 1%

Figure 2

Table 5. Level 2 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (nonrelease controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Target Dosage Form Weight
Filler	+/- 10%
Disintegrant	
starch	+/- 6%
other	+/- 2%
Binder	+/- 1%
Lubricant	
calcium or magnesium stearate	+/- 0.5%
other	+/- 2%
Glidant	
talc	+/- 2%
other	+/- 0.2%
Film coat	+/- 2%

Table 6. Level 3 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (nonrelease controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Target Dosage Form Weight	
Filler	Greater than +/- 10%	
Disintegrant		
starch	Greater than +/- 6%	
other	Greater than +/- 2%	
Binder	Greater than +/- 1%	
Lubricant		
calcium or magnesium stearate	Greater than +/- 0.5%	
other	Greater than +/- 2%	
Glidant		
talc	Greater than +/- 2%	
other	Greater than +/- 0.2%	
Film coat	Greater than +/- 2%	

Figure 3

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Table 7. Level 1 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (release controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Release	
	Controlling Excipient Content in the Modified	
	Release Solid Oral Dosage Form	
Any release controlling excipient(s)	+/- 5%	

Table 8. Level 2 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (release controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Releas	
	Controlling Excipient Content in the Modified	
	Release Solid Oral Dosage Form	
Any release controlling excipient(s)	+/- 10%	

Table 9. Level 3 Component and Composition Changes for Modified Release Oral Solid Dosage Forms (release controlling excipient)

Excipient	Percent Excipient (w/w) Out of Total Release	
	Controlling Excipient Content in the Modified	
	Release Solid Oral Dosage Form	
Any release controlling excipient(s)	Greater than +/- 10%	

Figure 4

Manufacturer Spectral Scan Spectral Data Sent to Data Base Packager or Distributor Communication with Database Verify Product in Secondary Inspection Product Library Library Maintained by Manufacturer or Third Party Vendor Pharmacy or Hospital Communication with Database Verify Secondary Product in Inspection Library Yes Patient

Table 10. Schematic of use within the commercial pipeline

Figure 5

Table 11. Composition of Aspirin Formulations

Component	Formulation A1 (mg/tab)	Formulation A2 (mg/tab)	Formulation A3 (mg/tab)
Aspirin	325	325	325
Microcrystalline cellulose	73	83	63
Magnesium stearate	2	2	2
TOTAL WEIGHT	400	410	390

Table 12. Composition of Prednisone Formulations

Component	Formulation B1 (mg/tab)	Formulation B2 (mg/tab)	Formulation B3 (mg/tab)
Prednisone	5	5	5
Microcrystalline cellulose	94.5	94.5	94.5
Magnesium stearate	0.5	0.75	0.25
TOTAL WEIGHT	100	100.25	99.75

Table 13. Composition of Indomethacin Formulations

Component	Formulation C1 (mg/tab)	Formulation C2 (mg/tab)	Formulation C3 (mg/tab)
Indomethacin	25	25	25
Microcrystalline cellulose	71.5	74	69
Croscarmellose sodium	3	2	4
Magnesium stearate	0.5	0.5	0.5
TOTAL WEIGHT	100	101.5	98.5

Table 14. Compositions of Acyclovir Formulations

Component	Formulation D1 (mg/tab)	Formulation D2 (mg/tab)	Formulation D3 (mg/tab)
Acyclovir	200	200	200
Microcrystalline cellulose	113.26	120.26	106.26
Starch	35	27.99	41.99
Magnesium stearate	1.75	1.75	1.75
TOTAL WEIGHT	350	350	350

Figure 6

WO 2005/031302

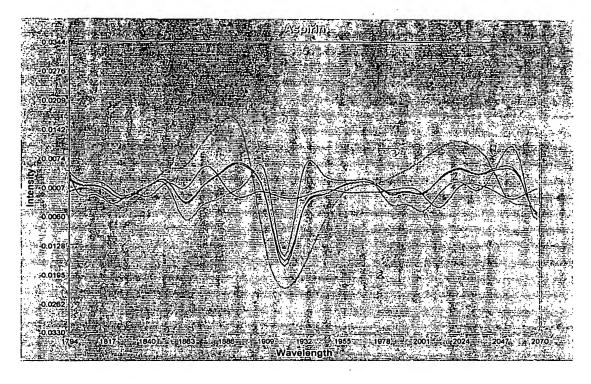


Table 15. 2nd Derivative of Absorbance vs. Wavelength: Aspirin Formulations

Figure 7

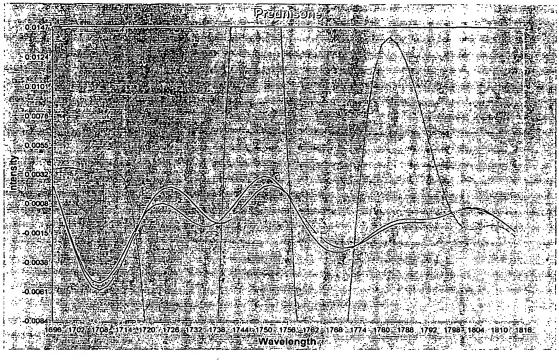


Table 16. 2nd Derivative of Absorbance vs. Wavelength: Prednisone Formulations

Figure 8

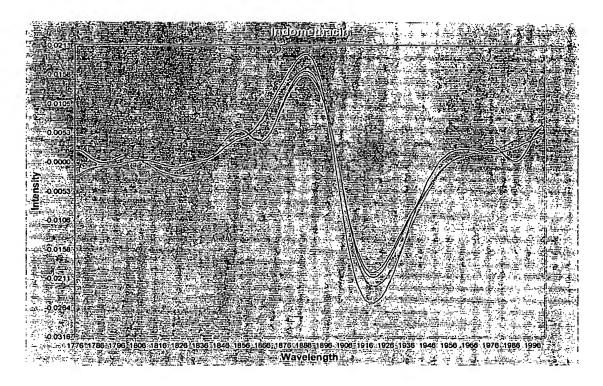


Table 17. $2^{\rm nd}$ Derivative of Absorbance vs. Wavelength: Indomethacin Formulations

Figure 9

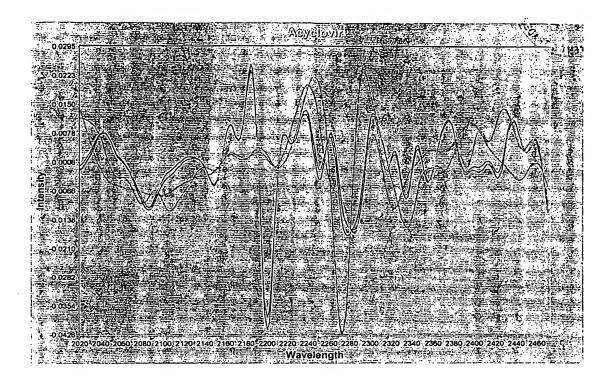


Table 18. 2nd Derivative of Absorbance vs. Wavelength: Acyclovir Formulations

Figure 10